

Serial No. 09/411,568
STN SEARCH

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=> help

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FILE COVERS 1967 - 12 Dec 2000 VOL 133 ISS 25
FILE LAST UPDATED: 11 Dec 2000 (20001211/ED)

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Serial No. 09/411,568
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=> s (itr or (internal (w) tandem (w) repeat#))/bi,ab

385 ITR/BI
319 ITR/AB

235311 INTERNAL/BI

209767 INTERNAL/AB

27102 TANDEM/BI

23200 TANDEM/AB

48590 REPEAT#/BI

44412 REPEAT#/AB

14 INTERNAL (W) TANDEM (W) REPEAT#

L1 398 (ITR OR (INTERNAL (W) TANDEM (W)

REPEAT#))/BI,AB

=> s ((anti (w) sense) or antisense)/bi,ab

223038 ANTI/BI

187535 ANTI/AB

20686 SENSE/BI

19836 SENSE/AB

849 ANTI (W) SENSE

17671 ANTISENSE/BI

14720 ANTISENSE/AB

L2 18196 ((ANTI (W) SENSE) OR ANTISENSE)/BI,AB

=> s 11 and 12

L3 5 L1 AND L2

=> s ribozyme#/bi,ab

4477 RIBOZYME#/BI

3235 RIBOZYME#/AB

L4 4477 RIBOZYME#/BI,AB

=> s 13 and 14

L5 2 L3 AND L4

=> d 13 1-5 bib ab

L3 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2000 ACS

AN 1999:795943 CAPLUS

DN 132:45813

TI Generation of recombinant adeno-associated virus vectors without formation of wild-type virus

IN Srivastava, Arun; Wang, Xu-Shan; Ponnazhagan, Selvarangan PA Advanced Research and Technology Institute, USA

SO PCT Int. Appl., 100 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9964569 A1 19991216 WO 1999-US13070 19990609 W:
AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH,
CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM,
HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK,
LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ,
PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR,
TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG,
KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE,
CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,
NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML,
MR, NE, SN, TD, TG

AU 9945587 A1 19991230 AU 1999-45587 19990609 PRAI US
1998-88714 19980610
WO 1999-US13070 19990609

AB A plasmid co-transfection system for the generation of recombinant adeno-assocd. virus 2 for use as a gene delivery virus that minimizes the generation of wild-type virus by preventing homologous recombination is described. Recombination is dependent upon 10 nucleotides of the viral D-sequence and helper vectors lacking sequence homol. in the D-sequence and helper plasmids lacking adenovirus inverted terminal repeats. Methods and compns. for the use of recombinant AAV plasmids and helper vectors lacking homol. in the D-sequence, and helper plasmids lacking the adenovirus ITRs for use in gene therapy are described. Mapping of recombination events leading to the generation of wild-type virus found most of them clustering in the 10 distal nucleotides of the D-sequence and also involved the inverted terminals repeats of the adenovirus 5 helper. Deletion of selected sequences gradually lowered the titer of wild-type virus to <0.1% of total virus.

RE.CNT 6

RE

(1) Qing; Journal of Virology 1998, V72(2), P1593
CAPLUS

(2) Wang; Journal of Molecular Biology 1995, V250, P573 CAPLUS (3) Wang; Journal of Virology 1996, V70(3), P1668 CAPLUS

(4) Wang; Journal of Virology 1997, V71(2), P1140
CAPLUS

(5) Wang; Journal of Virology 1997, V71(4), P3077
CAPLUS

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2000 ACS

AN 1999:244775 CAPLUS

DN 130:292438

TI Chimeric AAV/B19 parvovirus-based recombinant vector system specifically targeting the erythroid lineage

IN Srivastava, Arun; Ponnazhagan, Selvarangan PA Advanced Research and Technology Institute, USA
SO PCT Int. Appl., 76 pp.

CODEN: PIXXD2

DT Patent

Serial No. 09/411,568
STN SEARCH

LA English
FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9918227 A1 19990415 WO 1998-US21202 19981008 W:
AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN,
CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU,
ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA,
UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU,
TJ, TM RW: GE, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE,
CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,
NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML,
MR, NE, SN, TD, TG
AU 9912696 A1 19990427 AU 1999-12696 19981008 EP
1027451 A1 20000816 EP 1998-956097 19981008 R: AT, BE,
CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
PT, IE, FI
PRAI US 1997-61364 19971008
WO 1998-US21202 19981008

AB The present invention relates to the engineering, propagation and use of chimeric parvovirus vectors using sequences from adeno-assocd. virus (AAV) and B19 virus, which may be used to deliver genes to various target cells, including those of erythroid lineage. The system exploits the unique features of AAV and B19 such that it does not suffer from toxicity, oncogenicity, or immunogenicity concerns. Heterologous DNA sequences are cloned within the inverted terminal repeats (ITR) of AAV, without the presence of any AAV structural genes, and subsequently packaged inside the capsid structure of B19. Such a chimeric vector is achieved by creating a helper plasmid consisting of the rep gene of AAV, and the cap gene of B19. High titers of the vector may be generated, facilitating in vivo therapy. It is designed to specifically target primitive progenitor and differentiated cells of erythroid lineage, and can achieve stable integration and expression of transduced genes. RE.CNT 11
RE

(1) Childrens Hospital Inc; WO 9534670 A 1995 CAPLUS
(2) Latta, M; WO 9523867 A 1995 CAPLUS
(4) Ponnazhagan, S; Blood, Meeting Info: 39th Annual Meeting of the American Society of Hematology 1997 CAPLUS
(5) Ponnazhagan, S; J Virology 1998, V72(6), P5224 CAPLUS
(7) RES Corp Technologies Inc; WO 9309239 A 1993 CAPLUS
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2000 ACS
AN 1998:303188 CAPLUS
DN 129:77198
TI Site-specific integration in mammalian cells mediated by a new hybrid baculovirus-adeno-associated virus vector

AU Palombo, Fabio; Monciotti, Andrea; Recchia, Alessandra; Cortese, Riccardo; Ciliberto, Gennaro; La Monica, Nicola

CS IRBM P. Angeletti, Pomezia, 00040, Italy
SO J. Virol. (1998), 72(6), 5025-5034
CODEN: JOVIAM; ISSN: 0022-538X

PB American Society for Microbiology

DT Journal

LA English

AB Baculovirus can transiently transduce primary human and rat hepatocytes, as well as a subset of stable cell lines. To prolong transgene expression, we have developed new hybrid vectors which assoc. key elements from adeno-assocd. virus (AAV) with the elevated transducing capacity of baculovirus. The hybrid vectors contain a transgene cassette composed of the .beta.-galactosidase (.beta.-Gal) reporter gene and the hygromycin resistance (Hygr) gene flanked by the AAV inverted terminal repeats (ITRs), which are necessary for AAV replication and integration in the host genome. Constructs were derived both with and without the AAV rep gene under the p5 and p19 promoters cloned in different positions with respect to the baculovirus polyhedrin promoter. A high-titer prep. of baculovirus-AAV (Bac-AAV) chimeric virus contg. the ITR -Hygr-.beta.-Gal sequence was obtained with insect cells only when the rep gene was placed in an antisense orientation to the polyhedrin promoter. Infection of 293 cells with Bac-AAV virus expressing the rep gene results in a 10-to 50-fold increase in the no. of Hygr stable cell clones. Addnl., rep expression detd. the localization of the transgene cassette in the aav51 site in approx. 41% of cases as detected by both Southern blotting and fluorescent in situ hybridization anal. Moreover, site-specific integration of the ITR -flanked DNA was also detected by PCR amplification of the ITR -aav51 junction in transduced human fibro-blasts. These data indicate that Bac-AAV hybrid vectors can allow permanent, nontoxic gene delivery of DNA constructs for ex vivo treatment of primary human cells.

L3 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2000 ACS
AN 1998:169418 CAPLUS
DN 128:227084

TI Methods and compositions for liver-specific delivery of therapeutic molecules using recombinant adeno-associated virus vectors IN Srivastava, Aron; Ponnazhagan, Selvarangan; Chloemer, Robert H.; Wang, Xu-Shan; Yoder, Mervin C.; Zhou, Shang-Zhen; Escobedo, Jaime; Dwarki, Varavani
PA Chiron Corporation, USA; Indiana University
SO PCT Int. Appl., 32 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

Serial No. 09/411,568
STN SEARCH

PI WO 9809524 A1 19980312 WO 1997-US15453 19970902 W:
CA, JP
RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT,
LU, MC, NL, PT, SE EP 933997 A1 19990811 EP
1997-940762 19970902 R: AT, BE, CH, DE, DK, ES, FR,
GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI
PRAI US 1996-25616 19960906
US 1996-25649 19960911
WO 1997-US15453 19970902

AB Provided are methods for selectively expressing therapeutic mols., such as secretory proteins, antisense mols. and ribozymes, in the liver. The methods find use in treating hepatic diseases or conditions. The methods also find use in treating any disease or condition in which systemic administration of the therapeutic substance, for example, a secretory protein, is desired. The methods involve administering to a mammalian patient having a need for liver expression of a therapeutic mol. an AAV vector contg. a therapeutically effective amt. of the therapeutic mol. Also provided are novel vectors employable in these methods. Expts. revealed that, following i.v. injection of AAV vectors into mice, the AAV genomes were found predominantly in the liver. The heterologous genes carried by these vectors (chimeric cytomegalovirus promoter-lacZ or .beta.-globin promoter-globin genes) were expressed in the liver. Cotransfection of adenovirus 2-infected 293 cells with the AAV vectors and helper plasmid contg. cap and rep genes resulted in prodn. of 0.1-10% wild-type AAV. Replacement of the last 10 nucleotides of the ITR D sequence with unrelated nucleotides reduced this illegitimate recombination was reduced. Four recombinant AAV vectors (pD-5, pD-10, pD-15 and pD-20) with such modified ITR regions were prep'd.
L3 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2000 ACS
AN 1995:951301 CAPLUS
DN 123:332111
TI Integrative adenovirus expression vectors for use in gene therapy IN Denefle, Patrice; Latta, Martine; Perricaudet, Michel; Vigne, Emmanuelle PA
Rhone-Poulenc Rorer S.A., Fr.
SO PCT Int. Appl., 49 pp.

CODEN: PIXKD2

DT Patent

LA French

PAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 9523867 A1 19950908 WO 1995-FR233 19950228 W:
AM, AU, BB, BG, BR, BY, CA, CN, CZ, EE, FI, GE, HU,
JP, KE, KG, KP, KR, KZ, LK, LR, LT, LV, MD, MG, MN,
MW, MX, NO, NZ, PL, RO, RU, SD, SI, SK, TJ, TT, UA,
US, UZ, VN
RW: KE, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR,
GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG,
CI, CM, GA, GN, ML, MR, NE, SN, TD, TG
FR 2716893 A1 19950908 FR 1994-2445 19940303 FR
2716893 B1 19960412

CA 2184113 AA 19950908 CA 1995-2184113 19950228 AU
9518526 A1 19950918 AU 1995-18526 19950228 EP 748385
A1 19961218 EP 1995-910605 19950228 R: AT, BE, CH, DE,
DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE JP
09509578 T2 19970930 JP 1995-522730 19950228 ZA
9501803 A 19960109 ZA 1995-1803 19950303 US 6033885 A
20000307 US 1996-702573 19960912 PRAI FR 1994-2445
19940303

WO 1995-FR233 19950228

AB Recombination-defective adenoviruses carrying a cassette that can be integrated into the genome of host cells are constructed for use in gene therapy. The cassette particularly contains at least one inverted terminal repeat (ITR) of an adeno-assocd. virus (AAV) and a therapeutic gene. The use of the AAV ITR directs integration to the same locus in all cases and minimizes possible complications from random integration. The construction of virus carrying the lacZ reporter gene or a human lipoprotein AI gene under control of viral (vesicular stomatitis or Rous sarcoma virus) promoters is described.

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FILE COVERS 1967 - 7 Dec 2000 VOL 133 ISS 25
FILE LAST UPDATED: 7 Dec 2000 (20001207/ED)

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=> s 15

374 ITR/BI
309 ITR/AB

Serial No. 09/411,568
STN SEARCH

228219 INTERNAL/BI
203130 INTERNAL/AB
25935 TANDEM/BI
22201 TANDEM/AB
46369 REPEAT#/BI
42301 REPEAT#/AB
13 INTERNAL (W) TANDEM (W) REPEAT#
215262 ANTI/BI
180573 ANTI/AB
19151 SENSE/BI
18427 SENSE/AB
813 ANTI (W) SENSE
16873 ANTISENSE/BI
14081 ANTISENSE/AB
4250 RIBOZYME#/BI
3054 RIBOZYME#/AB
L6 2 L3 AND L4

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TOTAL ENTRY SESSION CA SUBSCRIBER PRICE 0.00 -2.78
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